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2 1. A compound of the formula:

4 a pharmaceutically acceptable salt or a prodrug thereof,

5 wherein

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- R^1 is alkyl or $-NR^7R^8$, where each of R^7 and R^8 is independently
- 7 hydrogen or alkyl;
- 8 R² is hydrogen or alkyl;
- each of R³, R⁴, R⁵, and R⁶ is independently hydrogen, halide, alkyl,
- 10 –OR⁹ (where R⁹ is hydrogen, alkyl, a hydroxy protecting group,
- or cycloalkylalkyl), -SR¹⁰ (where R¹⁰ is hydrogen or alkyl), or
- 12 $-NR^{11}R^{12}$ (where each of R^{11} and R^{12} is independently
- hydrogen, alkyl, or a nitrogen protecting group), provided R³,
- 14 R⁴, R⁵, and R⁶ are not all simultaneously alkyl); or R³ and R⁴
- together with atoms to which they are attached to form
- heterocyclyl, heteroaryl, or cycloalkyl; and
- 17 R¹⁴ is hydrogen, lower alkyl or -OR¹⁵, where R¹⁵ is hydrogen, lower
- alkyl, or a hydroxy protecting group.
- 1 2. The compound according to Claim 1, wherein R¹⁴ is hydrogen.
- 1 3. The compound according to Claim 2, wherein R¹ is alkyl.
- 1 4. The compound according to Claim 3, wherein R¹ is selected from the 2 group consisting of methyl, ethyl, and isopropyl.
- 1 5. The compound according to Claim 3, wherein R² is hydrogen.
- 1 6. The compound according to Claim 5, wherein each of R⁷ and R⁸ is independently hydrogen or methyl.

1	7.	The compound according to Claim 6, wherein each of R ³ , R ⁴ , R ⁵ , and	
2	R ⁶ is independently hydrogen, halide, alkyl, or -OR ⁹ , where R ⁹ is hydrogen, alkyl, a hydroxy		
3	protecting group, or cycloalkylalkyl; or R ³ and R ⁴ together with atoms to which they are		
4	attached to form heterocyclyl, heteroaryl, or cycloalkyl.		
1	8.	The compound according to Claim 7, wherein at least one of R ³ , R ⁴ ,	
2	R ⁵ , and R ⁶ is alkyl, halide, or -OR ⁹ , where R ⁹ is as defined in Claim 1.		
1	9.	The compound according to Claim 8, wherein at least one of R ³ , R ⁴ ,	
2	R ⁵ , and R ⁶ is bromo,	chloro, fluoro, methoxy, ethoxy, methyl, and hydroxy.	
1	10.	The compound according to Claim 9, wherein	
2	(a)	R ³ is methoxy, and R ⁴ , R ⁵ , and R ⁶ are hydrogen;	
3	(b)	R ³ is methyl, R ⁶ is methoxy, and R ⁴ and R ⁵ are hydrogen;	
4	(c)	R ³ is methyl, R ⁶ is chloro, and R ⁴ and R ⁵ are hydrogen;	
5	(d)	R ³ is chloro, R ⁴ is methoxy, and R ⁵ and R ⁶ are hydrogen;	
6	(e)	R ³ is methyl, R ⁴ is chloro, and R ⁵ and R ⁶ are hydrogen;	
7.	- (f)	R ³ is methyl, R ⁴ is methoxy, and R ⁵ and R ⁶ are hydrogen;	
8	(g)	R^4 is chloro, and R^3 , R^5 and R^6 are hydrogen;	
9	(h)	R^4 is methoxy, and R^3 , R^5 , and R^6 are hydrogen.	
10	(i)	R ³ is methyl, R ⁶ is bromo, and R ⁴ and R ⁵ are hydrogen;	
11	(j)	R ³ is bromo, R ⁴ is methoxy, and R ⁵ and R ⁶ are hydrogen;	
12	(k)	R ³ is methyl, R ⁴ is bromo, and R ⁵ and R ⁶ are hydrogen;	
13	(1)	R ⁴ is bromo, and R ³ , R ⁵ and R ⁶ are hydrogen; or	
14	(m)	R ³ is ethoxy and R ⁴ , R ⁵ and R ⁶ are hydrogen.	
1	11.	The compound according to Claim 7, wherein R ³ and R ⁴ together with	
2	atoms to which they are attached to form furanyl, dihydrofuranyl, or pyrrolyl.		

1 12. The compound according to Claim 11, wherein R³ and R⁴ together 2 with atoms to which they are attached to form furanyl or dihydrofuranyl.

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2 the formula:

4

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- 1 14. A method for producing an imidazolin-2-ylmethyl-substituted aromatic
- 2 compound of the formula:

$$\mathbb{R}^{1}$$

$$\mathbb{R}^{2}$$

$$\mathbb{R}^{2}$$

$$\mathbb{R}^{6}$$

$$\mathbb{R}^{14}$$

$$\mathbb{R}^{14}$$

$$\mathbb{R}^{14}$$

3

4 said method comprising contacting a nitrile compound of the formula:

5 6

- with ethylene diamine to produce the imidazolin-2-ylmethyl-substituted aromatic compound,
- 7 wherein
- R¹ is alkyl, -NR⁷R⁸, where each of R⁷ and R⁸ is independently hydrogen or alkyl;
- 10 R² is hydrogen or alkyl;
- each of R³, R⁴, R⁵, and R⁶ is independently hydrogen, halide, alkyl, -OR⁹,
- where R⁹ is hydrogen, alkyl, a hydroxy protecting group, or
- cycloalkylalkyl, –SR¹⁰, where R¹⁰ is hydrogen or alkyl, or –NR¹¹R¹²,
- where each of R¹¹ and R¹² is independently hydrogen, alkyl, or a
- nitrogen protecting group, provided R³, R⁴, R⁵, and R⁶ are not all
- simultaneously alkyl); or R³ and R⁴ together with atoms to which they
- are attached to form heterocyclyl, heteroaryl, or cycloalkyl; and
- 18 R¹⁴ is hydrogen, lower alkyl or -OR¹⁵, where R¹⁵ is hydrogen, lower alkyl, or
- a hydroxy protecting group.

2 compound of the formula:

4 said method comprising contacting an ester compound of the formula:

$$R^{4}$$

$$R^{2}$$

$$R^{2}$$

$$R^{6}$$

$$R^{14}$$

$$R^{14}$$

$$R^{14}$$

$$R^{13}$$

5 6

7

8

1

2

1

2

1

3

with ethylene diamine in the presence of a trialkylaluminum to produce the imidazolin-2-

ylmethyl-substituted aromatic compound,

wherein

R¹ is alkyl, -NR⁷R⁸, where each of R⁷ and R⁸ is independently hydrogen or 9 alkyl; 10

R² is hydrogen or alkyl; 11

each of R³, R⁴, R⁵, and R⁶ is independently hydrogen, halide, alkyl, -OR⁹, 12 where R⁹ is hydrogen, alkyl, a hydroxy protecting group, or 13 cycloalkylalkyl, -SR¹⁰, where R¹⁰ is hydrogen or alkyl, or -NR¹¹R¹², 14 where each of R¹¹ and R¹² is independently hydrogen, alkyl, or a 15 nitrogen protecting group; or R³ and R⁴ together with atoms to which 16

they are attached to form heterocyclyl, heteroaryl, or cycloalkyl; 17

R¹³ is alkyl; and 18

R¹⁴ is hydrogen, lower alkyl or -OR¹⁵, where R¹⁵ is hydrogen, lower alkyl, or 19 a hydroxy protecting group. 20

16. The method of Claim 15, wherein the trialkylaluminum is trimethylaluminum or triethylaluminum.

17. A composition comprising:

- a therapeutically effective amount of a compound of Claim 1; and (a)
- 3 (b) a pharmaceutically acceptable carrier.

1	18.	A method for treating a disease state selected from the groups	
2	consisting of urge incontinence, stress incontinence, overflow incontinence, functional		
3	incontinence, sexual dysfunction, nasal congestion, and CNS disorders selected from the		
4	group depression, anxiety, dementia, senility, Alzheimer's, deficiencies in attentiveness and		
5	cognition, eating disorders, obesity, bulimia and anorexia, said method comprising		
6	administering to a patient in need of such treatment a therapeutically effective amount of a		
7	compound of Claim 1.		
1	19.	A method for treating a disease state comprising urinary incontinence	
2	by administering to a subject in need of such treatment an effective amount of a Compound		
3	of Claim 1.		
4			
1	20.	The method of Claim 19, wherein the disorder is stress incontinence.	
1	21.	The method of Claim 19, wherein the disorder is urge incontinence.	
1	22.	A method for treating nasal congestion by administering to a mammal	
2	in need of such treatment an effective amount of a Compound of Claim 1.		
1	23.	The method of Claim 22, wherein the disorder is nasal congestion.	
1	24.	The method of Claim 23, wherein the disorder is sinusitis or otitis.	
1	24.	The medica of claim 23, wherein the disorder is smashes of others.	
1	25.	A method for treating sexual dysfunction by administering to a	
2	mammal in need of such treatment an effective amount of a Compound of Claim 1.		